

CLAIMS

We claim:

1. A composition of matter comprising an isolated polypeptide selected from the group consisting of SEQ ID NO:1, SEQ ID NO:2, SEQ ID NO:3, SEQ ID NO:4, SEQ ID NO:5, SEQ ID NO:6, SEQ ID NO:7 and variants thereof.
2. The composition of claim 1, further comprising a carrier.
3. A method of detecting presence of antibodies to *Ehrlichia* comprising:
 - (a) contacting one or more polypeptides selected from the group consisting of the polypeptides shown in SEQ ID NO:1, SEQ ID NO:2, SEQ ID NO:3, SEQ ID NO:4, SEQ ID NO:5, SEQ ID NO:6, SEQ ID NO:7, and variants thereof, with a test sample suspected of comprising antibodies to *Ehrlichia*, under conditions that allow polypeptide/antibody complexes to form;
 - (b) detecting polypeptide/antibody complexes;
wherein the detection of polypeptide/antibody complexes is an indication that antibodies to *Ehrlichia* are present in the test sample.
4. The method of claim 3, further comprising contacting the complexes of step (a) with an indicator reagent comprising a signal generating compound that generates a measurable signal prior to the performance of step (b).
5. The method of claim 3, wherein the presence of antibodies to *Ehrlichia canis* are detected.

6. The method of claim 3, wherein the presence of antibodies to *Ehrlichia chaffeensis* are detected.

7. The method of claim 3, wherein the antibodies are fragments of antibodies.

8. The method of claim 3 wherein the amount of antibody in a test sample is determined.

9. The method of claim 3, wherein the polypeptide is attached to a substrate.

10. The method of claim 3, wherein the polypeptide provided is shown in SEQ ID NO:1.

11. The method of claim 3, wherein the polypeptide provided is shown in SEQ ID NO:2.

12. The method of claim 3, wherein the polypeptide provided is shown in SEQ ID NO:3.

13. The method of claim 3, wherein the polypeptide provided is shown in SEQ ID NO:4.

14. The method of claim 3, wherein the polypeptide provided is shown in SEQ ID NO:5.

15. The method of claim 3, wherein the polypeptide provided is shown in SEQ ID NO:6.

16. The method of claim 3, wherein the polypeptide provided is shown in SEQ ID NO:7.

17. The method of claim 3, wherein the one or more polypeptides are provided in a multimeric form.

18. The method of claim 3, wherein the test sample is a biological sample obtained from a mammal.

19. The method of claim 18, wherein the mammal is selected from the group consisting of humans and dogs.

20. The method of claim 3 wherein the method comprises an assay selected from the group of assays consisting of a reversible flow chromatographic binding assay, an enzyme linked immunosorbent assay, a western blot assay, and an indirect immunofluorescence assay.

21. A device containing one or more polypeptides selected from the group consisting of the polypeptides shown in SEQ ID NO:1, SEQ ID NO:2, SEQ ID NO:3, SEQ ID NO:4, SEQ ID NO:5, SEQ ID NO:6, SEQ ID NO:7, and variants thereof.

22. The device of claim 21, further comprising instructions for use of the one or more polypeptides for the identification of an *Ehrlichia* infection in a mammal.

23. The device of claim 22, wherein the identification of an *Ehrlichia* infection is done using a method of detecting presence of antibodies to *Ehrlichia* comprising:

- (a) contacting one or more polypeptides selected from the group consisting of the polypeptides shown in SEQ ID NO:1, SEQ ID NO:2, SEQ ID NO:3, SEQ ID NO:4, SEQ ID NO:5, SEQ ID NO:6, SEQ ID NO:7, and variants thereof, with a test sample suspected of comprising antibodies to *Ehrlichia*, under conditions that allow polypeptide/antibody complexes to form;
- (b) detecting polypeptide/antibody complexes;

wherein the detection of polypeptide/antibody complexes is an indication that an *Ehrlichia* infection is present.

24. The device of claim 22, wherein the *Ehrlichia* infection is caused by *Ehrlichia canis* or *Ehrlichia chaffeensis*.

25. An article of manufacture comprising packaging material and, contained within the packaging material, one or more polypeptides selected from the group consisting of the polypeptides shown in SEQ ID NO:1, SEQ ID NO:2, SEQ ID NO:3, SEQ ID NO:4, SEQ ID NO:5, SEQ ID NO:6, SEQ ID NO:7, and variants thereof.

26. The article of manufacture of claim 25 wherein the packaging material comprises a label that indicates that the one or more polypeptides can be used for the identification of *Ehrlichia* infection in a mammal.

27. The article of manufacture of claim 26, wherein the identification of an *Ehrlichia* infection is done using a method of detecting presence of antibodies to *Ehrlichia* comprising:

- (a) contacting one or more polypeptides selected from the group consisting of the polypeptides shown in SEQ ID NO:1, SEQ ID NO:2, SEQ ID NO:3, SEQ ID NO:4, SEQ ID NO:5, SEQ ID NO:6, SEQ ID NO:7, and variants thereof, with a test sample suspected of comprising antibodies to *Ehrlichia*, under conditions that allow polypeptide/antibody complexes to form;
- (b) detecting polypeptide/antibody complexes;

wherein the detection of polypeptide/antibody complexes is an indication that an *Ehrlichia* infection is present.

28. The article of manufacture of claim 26, wherein the *Ehrlichia* infection is caused by *Ehrlichia canis* or *Ehrlichia chaffeensis*.

29. A method of diagnosing an *Ehrlichia* infection in a mammal comprising:

(a) obtaining a biological sample from a mammal suspected of having an *Ehrlichia* infection;

(b) contacting one or more polypeptides selected from the group consisting of the polypeptides shown in SEQ ID NO:1, SEQ ID NO:2, SEQ ID NO:3, SEQ ID NO:4, SEQ ID NO:5, SEQ ID NO:6, SEQ ID NO:7, and variants thereof, with the biological sample under conditions that allow polypeptide/antibody complexes to form;

(c) detecting polypeptide/antibody complexes;

wherein the detection of polypeptide/antibody complexes is an indication that the mammal has an *Ehrlichia* infection.

30. The method of claim 29 further comprising contacting the complexes of step (b) with an indicator reagent comprising a signal generating compound that generates a measurable signal prior to the performance of step (c).

31. The method of claim 29, wherein the *Ehrlichia* infection is caused by *Ehrlichia canis*.

32. The method of claim 29, wherein the *Ehrlichia* infection is caused by *Ehrlichia chaffeensis*.

33. The method of claim 29, wherein the mammal is a human or a dog.

34. A monoclonal antibody that specifically binds to at least one epitope of an *Ehrlichia canis* or *Ehrlichia chaffeensis* polypeptide, said polypeptide selected from the group consisting of SEQ ID NO:1, SEQ ID NO:2, SEQ ID NO:3, SEQ ID NO:4, SEQ ID NO:5, SEQ ID NO:6, and SEQ ID NO:7.